



Message from VPRA, Dr. Robert Nobles



Hello from the Office of Research Administration! Great things are happening in ORA and we are excited to share with you some of the initiatives, strategic actions, and plans for the 2025 fiscal year.

Please take a few minutes to watch this special video message from Dr. Robert Nobles. As always, thank you for your ongoing commitment to research excellence and partnership!

Mark Your Calendar! Research Administration Townhall Meeting with the VPRA

Research Administration Townhall Meeting with the VPRA



Robert Nobles, DrPH, MPH
Vice President
Research Administration



Kim Maune, MHA
Chief Implementation Officer, GMS
Executive Associate Dean of
Administration & Finance
Chief Business Officer
Rollins School of Public Health

Wednesday, October 23, 2024

12:00 PM – 1:00 PM ET

Event Description: The Office for Research Administration's VPRA is hosting Research Administration's 2024 Townhall meeting. This meeting will be split to share ORA Updates (FY24 progress and FY25 strategic focus) and a Grants Management System (GMS) Update and Demo.

Advance Questions: You may post questions in advance via [Qualtrics](#) until 4 PM, **Tuesday, October 22nd**.

We will address submitted questions during the meeting. You will also have a chance to submit questions during the live meeting.

Audience: University-wide

[Register Here](#)

[Submit a Question](#)

Grants Management System Update: Implementing Insight



Author: Kim Maune, Emory Chief Implementation Officer, Insight GMS

Emory University is investing in a **fully integrated research administration grants management package**. The product, Insight, is a cloud-based system unifying distributed and complex organizational processes into one highly integrated system to enable compliance, efficiency, and transparency.

Our research systems are currently numerous and not integrated, creating duplicate entry and inefficient workflow, increasing administrative burden. Implementation and optimization of a GMS will standardize research processes across Emory to support faculty, research staff, study teams, and administrative offices to manage their research portfolios efficiently and effectively.

[Read Full Article](#)

FY24 Q1 - Q3 Mission Metrics Now Available

Author: Alex Wagner, Director, Research Data Analytics, (RBO)

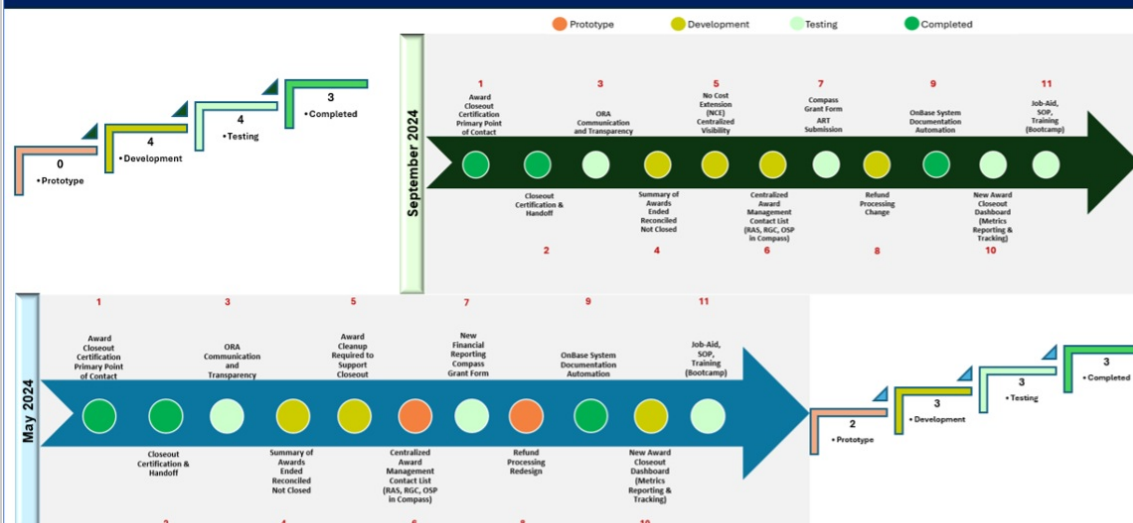
[View the FY24 Mission Metrics Report](#)

What are Mission Metrics?: This dashboard was developed as a reporting process for faculty facing metrics. The goal is to supply meaningful metrics to faculty to measure performance of ORA's departments and units. This data-informed dashboard will seek to improve ORA's transparency and identify areas for improvement. The metrics are produced in several phases to incorporate feedback and develop additional metrics.

The FY24 and Past Mission Metrics are now available on the [RBO website](#).

Emory's Award Closeout Plan and Roadmap

Award Closeout Plan & Roadmap Progress Update



Author: Patrick Amihire, Associate Director (RGC)

This month's article is focused on the progress of **Emory's Award Closeout Processes/Projects Development** and a **Summary of the Revised OMB guidance also known as the Uniform Guidance (UG)**. First, the recent changes to the federal guidelines provide additional reasons why our institution must reduce the administrative burden relative to closing awards effectively and timely to be compliant with Sponsoring Agencies' terms and conditions. Secondly, the ongoing initiatives related to Award Closeout Processes/Projects are progressing steadily to facilitate compliance with both UG and Emory policies and guidance.

[Read Full Article](#)

FY2024 Research Administration Annual Satisfaction Survey

Dear Colleagues,

The Office of Research Administration has launched our annual researcher survey

to solicit your feedback on the progress that is being made to support the excellence within our research enterprise.

In our efforts to build the most responsive research administration for Emory, we would appreciate your feedback on your experiences over the past year. Your feedback will enable us to align research administrative services with your needs. **This annual survey is anonymous.**

Your feedback will help enhance our understanding of current needs, while helping us document where critical progress has been made. We utilized past year's feedback to make improvements in several areas, including moving forward with modernizing our IT systems, increasing integration, and delivering transparency.

FY2024 Annual Satisfaction Survey

Please note: This survey is not intended for Research Administration staff

The survey should require no more than 5-10 minutes of your time, with the option to provide more thorough commentary as you wish.

The survey will be active until **COB October 3rd, 2024** and this year's results will again be shared in various formats and meetings **beginning in 2025**. Thank you for your time and attention. We again look forward to hearing and implementing your feedback!

Thank you in advance for your partnership,

Robert

Robert Nobles, DrPH, MPH
Vice President for Research Administration

[Coming in May 2025: Investigators Must Use SciENCv To Format NIH and NSF Personnel Documents](#)



Authors: Deepika Bhatia, AVP, (RCRA) & Kimberly Eck, Sr. AVP, (SVPR)

SciENCv is an application in My NCBI that helps you create and manage documents in support of grant applications with participating agencies. In SciENCv you can document your education, employment, research activities, publications, honors, research grants, and other professional contributions.

<https://www.ncbi.nlm.nih.gov/sciencv/>

For NIH: Biosketches and Current and Pending Support for proposals and RPPRs submitted **on or after May 25, 2025** must use the new Common Form and be produced using SciENCv. NSF has already implemented the use of SciENCv. Please see [NOT-OD-24-163](#) for details.

[Read Full Article](#)

[Press Release: PRIM&R Releases Podcast to Explore Scientific Research With Public](#)



Research Ethics Reimagined
PRIM&R PUBLIC RESPONSIBILITY IN MEDICINE AND RESEARCH



BOSTON, September 12, 2024 /PRIM&R/ – Public Responsibility in Medicine and Research (PRIM&R) announced the release of its podcast, “Research Ethics Reimagined,” designed to help further the public’s understanding of scientific research.

The monthly podcast, hosted by PRIM&R’s Executive Director Ivy R. Tillman, EdD, CIP, features conversations with “scientists, researchers, bioethicists and some of the leading minds exploring new frontiers of science,” according to the official show summary.

“From teaching to navigating the complexities of research ethics as an IRB director, my career has been shaped by a commitment to bridging gaps, fostering understanding, and helping others make connections,” Tillman said of her role as the podcast’s host.

PRIM&R has released four podcast episodes to date:

- [Episode 1](#) (“Making Science More Accessible”) focuses on building and maintaining trust in research and science in the 21st century by creating a connection with the public and simplifying the language scientists use.

Guests: **Robert Nobles**, DrPH, MPH, CIP, Vice President for Research Administration at Emory University, and **Amanda M. Dettmer**, PhD, a research scientist at the Yale Child Study Center.

[Read Full Article](#)

Federal Agency Updates

NIH Automated Email Seeking Just-in-Time Information to Be Retired October 1



An automated email from eRA for Just-in-Time information, sent for applications with an overall impact score of 30 or less, will be retired on October 1, 2024.

[Read Full Article](#)

Changes to Data Management and Sharing (DMS) Plan Progress Reporting and the Submission of Revised DMS Plans Are Coming on October 1



On October 1, NIH is adding several new Data Management and Sharing (DMS) questions to Research Performance Progress Reports (RPPRs) and updating the process for submitting revised DMS Plans to NIH for review.

[Read Full Article](#)

Recording: Revision of NSF Award Terms and Conditions Implementing Revised 2 CFR



The NSF Policy Office in the Division of Institution and Award Support (DIAS) held a webinar on September 12 at 2:00 PM ET which provided the framework for these revisions and how they affect NSF proposals and awards.

[View Recording](#)

[Learn More](#)

Summary: NIH Closeout Process Webinar



Summary: NIH Closeout Process Webinar



Closeout Roles and Responsibilities

Recipient	Subrecipient	NIH Staff
<ul style="list-style-type: none">• Primary compliance with all terms and conditions of the Notice of Award.• Ensure subrecipient compliance with all terms and conditions of the award.• Submission of timely and accurate closeout reports.• Timely communication with NIH staff.	<ul style="list-style-type: none">• The terms and conditions flow down to subrecipients in accordance with 2 CFR Part 200.101(b)(2) – this includes closeout requirements.• Compliance with applicable terms and conditions of the award and the written consortium agreement.• Timely communication and engagement with the recipient.	<ul style="list-style-type: none">• Review and acceptance/approval of closeout reports.• Ensuring closeout within one year from the project period end date.• Initiate unilateral closeout if recipient does not provide timely accurate reports.• Report all unilaterally closed awards for noncompliance in SAM.gov Responsibility/Qualification.

Authors: Tricia Callahan, Interim Director, (OSOT) & Edwin Bommel, AVP, (RGC)

The webinar summary link below reviews the NIH Closeout Process webinar presented by the NIH Office of Extramural Research (OER) on September 17, 2024. The webinar recording, transcript, and presentation slides can be found on the NIH Event Page.

[Webinar Summary](#)

[Presentation Slides](#)

[NIH Event Page](#)

Research Administration Training Opportunities

ORA Boot Camp Series: NIH Unplugged



EMORY
UNIVERSITY

Office of Strategic
Optimization and Training

Author: Tricia Callahan, Interim Director, (OSOT)

The NIH Unplugged Series offers an in-depth exploration of the NIH grant application and management process. Designed for investigators and administrators alike, it provides essential knowledge and practical guidance for navigating the complexities of NIH funding.

Through informative presentations and interactive discussions, the NIH Unplugged Series equips participants with the tools and knowledge necessary to increase their chances of securing NIH funding and successfully managing awarded grants.

NIH Unplugged Webinar
List

Other Research Training
Up and Coming Events

Department Updates

Environmental Health and Safety Office

[H5 Avian Influenza Virus Exemption](#)

Until recently H5N1 Avian Influenza was regulated under the USDA/CDC Select Agent Program. During this temporary exemption Emory researchers will soon begin work with this virus in the highest level of containment at Emory including Biosafety Level 3 (BSL3) and Animal Biosafety Level 3 (ABSL3).



[Read Full Article](#)

Institutional Review Board



Planning to Submit a Federal Grant for Multi-Site Human Research?

Reach out to the Emory IRB reliance team at irb.reliance@emory.edu to discuss the single IRB plan before you submit!

Also, a reminder about our Collaborative Research webpage here, with instructions for all things related to inter-IRB reliance: <https://irb.emory.edu/guidance/research-types/collaborative.html>

Collaborative Research/Single IRBs/Reliance Agreements

Collaborative Research

Collaborative research is research that may be conducted by researchers from more than one institution at just one site or when researchers from multiple institutions conduct different parts of a protocol. For example, non-Emory investigators may assist the Emory study team in conducting a research study at Emory. Another example is when an Emory study team conducts research interventions but another institution conducts the data analysis.

A critical first step is to determine whether Emory or the external collaborators are considered "engaged" in human subjects research. We have created the [Engagement Guidance Checklist](#) to assist you in making this determination. Once you complete the checklist, upload it to the protocol section of the smart form. Reliance agreements are not required for sites and/or external collaborators when an Emory researcher is not engaged in human subjects research.

If you determine the external collaborator is engaged in human subjects research, you may add external collaborators to an Emory IRB approved study if the research is federally funded and the research was not determined to be exempt. There are other scenarios where we may agree to provide oversight for external collaborators. Reach out to the reliance team at irb.reliance@emory.edu to determine if you can add the external collaborator(s).

If you want to add an external collaborator to an Emory IRB approved study, follow the steps below:

- Send an email to irb.reliance@emory.edu with the following information:
 - name of the external study team member(s) you wish to add to your submission
 - name of the external study team member's home institution
 - description of the specific research activities the external study team member will perform
 - time period the external team member is expected to conduct research activities
 - a link to the study in eIRB
 - the completed [External Study Team Member List Template](#) (use only one list per study and include all external study team members)

Effective October 1: Protocol and Consent Template Updates

The IRB has released new [protocol](#) and [consent](#) templates for biomedical research, available on the IRB website. Additional templates will be updated in the near future.

[Read Full Article](#)



Forms and Templates

WAIVERS

Find the waivers you need or browse FAQs created to help you use them.

[VIEW WAIVERS](#)

CONSENT TOOLKIT

Learn how and when to use consent protocol and documentation.

[VIEW CONSENT TOOLKIT](#)

FOOD AND DRUG ADMINISTRATION (FDA)

Browse forms and templates created to help with your FDA-influenced projects.

[VIEW FDA RESOURCES](#)

OTHER

Find other forms and templates made to help you navigate your research projects.

[BROWSE OTHER RESOURCES](#)



New Members Sought for IRB Service

The IRB is seeking new members especially from the following areas: AI/ML, Anthropology, Emergency Medicine, Endocrinology, Infectious Disease, OB/GYN, Pediatrics, Pulmonology, Psychiatry, Sociology, and Surgery/Transplant. Faculty from other areas are also welcome to inquire. Please reach out to Carol Corkran and Rebecca Rousselle, at ccorkra@emory.edu and rrouss2@emory.edu. You can find information about membership here: <https://irb.emory.edu/members/index.html>

Members

- [Expectations for Committee Members \(PDF\)](#)
- A training program is required of prospective members to ensure that they have a solid foundation in theory, regulations, and practical skills before engaging in service with the Emory IRB.
- The new-member orientation includes the following: observation of an Emory IRB meeting; participation in a training session on IRB Policies and Procedures; completion of the CITI online training program (if not currently certified); and attendance at a hands-on eIRB training session.
- Once the training steps noted above are complete, the Vice President for Research Administration at Emory will sign an appointment letter.
- Members are encouraged to contribute comments to the discussions and to demonstrate respect for each other's opinions
- Members are expected to review all applications on the agenda, whether assigned as primary/secondary reviewer or not.
- For more information on becoming an Emory IRB member, please contact [Rebecca Rousselle](mailto:Rebecca.Rousselle).

Office of Research Development

[Grant Forward at Emory University](#)

The Office of the Senior Vice President for Research sponsors a university-wide subscription to [GrantForward](#).

GrantForward provides easy, customizable access to thousands of grant opportunities through a single easy-to-use interface. Researchers can use this service to self-curate personalized funding notices, as well as to create research profiles that will further refine and individualize grant recommendations provided by the system.

[Read Full Article](#)



[Effective January 25, 2025 – A Revised NIH Grant Review Process with a Focus on Scientific Merit & Reputational Bias Reduction](#)

The National Institutes of Health (NIH) has taken steps to simplify its grant review process for most research project grants for applications with due dates of January 25, 2025 or later. Upcoming changes have been undertaken in order to address the complexity of the peer review process and the potential for reputational bias to affect peer review outcomes.

[Read Full Article](#)

Research Compliance & Regulatory Affairs

Emory IACUC September Newsletter

Check out the IACUC September 2024 newsletter for updates on the following topics:

- IACUC Policies
- Post Approval Attestation of Responsibilities
- From the Office of Occupational Health and Safety
- DEA warning for DAR Vets and Investigators
- Alerts to Researchers
- Alternative Searches in Protocol
- Ask RCRA – September 19, 2024, 12 noon
- Have you read the most recent version of your approved IACUC protocol?
- Resources for researchers- What is continuing education?
- IACUC Site Inspections – New schedule for the second period of 2024 6/1/2024- 11/30/2024
- IACUC Office Contacts

[Read the September 2024 Issue](#)

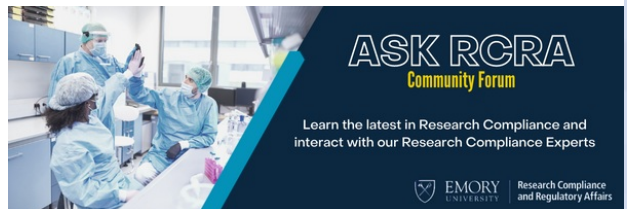
October "Ask RCRA" Community Forum

The Office of Research Compliance and Regulatory Affairs (RCRA) is delighted to invite you to our monthly "Ask RCRA" community forum! Please join us for important research compliance information, including regulatory updates and institutional changes in policies and procedures.

*** Take the survey and let us know about research compliance topics you'd like to see when we return!

- [Ask RCRA Feedback Survey](#) ***

Do you have any questions we can answer during the session? Please submit questions to: researchcompliance@emory.edu



When: Thursday, October 17, 2024
Time: 12PM - 1PM ET

[Register Here](#)

RCRA Update Highlights

[NSPM 33](#)

[Policy 7.21 Minors Participating in Research Activities at Emory](#)

The federal government has updated the standards surrounding disclosure, including requirements and processes. For additional information, please see the [OSTP Guide](#).

New revisions to this policy clarify the definition of a minor and requires PIs/Sponsors who will host minors in their lab to complete a Minor Registration Form. This form can be obtained from by emailing the research compliance office: researchcompliance@emory.edu.

iThenticate

A new plagiarism detection software is being [implemented at Emory](#). This software is a tool to ensure research documents including manuscripts and grant applications follow proper citation practices. This system currently in use at other major Universities across the country including Harvard, Stanford and UC Berkley.

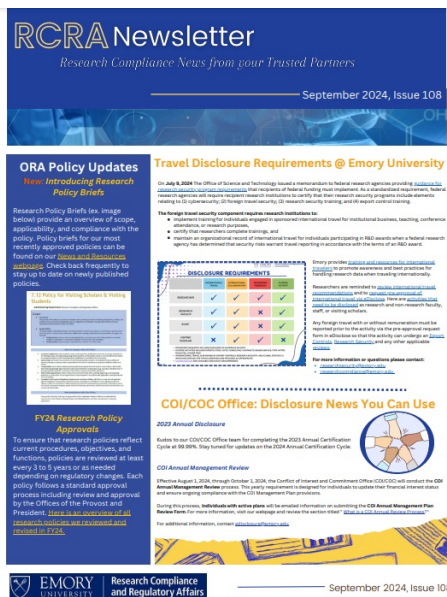
Controlled Substances & Dangerous Drugs

Ensuring compliance with regulations governing controlled substances and dangerous drugs is paramount in research. Here are important pointers and resources to support your use of controlled substances in research -<https://rcra.emory.edu/includes/documents/sections/oric/cs-dd-one-pager-pi.pdf>

Emory RCRA September Newsletter

Check out the RCRA September 2024 newsletter for updates on the following topics:

- ORA Policy Updates
- Travel Disclosure Requirements @Emory University
- COI/COC Office: Disclosure News You Can Use
- Use of Controlled Substances and Dangerous Drugs in Research
- Emory IACUC September Newsletter
- Minors Participating in Research Activities @Emory



Read the September 2024 Issue

Faculty Feedback Form

Do you have feedback in reference with Research Administration? We would love to hear from you.

Submit Faculty Feedback



View this newsletter and past editions on our [ORA Newsletter webpage](#)